LA-UR-01-6653

Approved for public release; distribution is unlimited.

Title:	SCORR - Supercritical Carbon Dioxide Resist Removal
Author(s):	Gunilla B. Jacobson , Laurie Williams, W. Kirk Hollis Jerome Barton and Craig M. V. Taylor
Submitted to:	http://lib-www.lanl.gov/la-pubs/00796632.pdf

Los Alamos National Laboratory, an affirmative action/equal opportunity employer, is operated by the University of California for the U.S. Department of Energy under contract W-7405-ENG-36. By acceptance of this article, the publisher recognizes that the U.S. Government retains a nonexclusive, royalty-free license to publish or reproduce the published form of this contribution, or to allow others to do so, for U.S. Government purposes. Los Alamos National Laboratory requests that the publisher identify this article as work performed under the auspices of the U.S. Department of Energy. Los Alamos National Laboratory strongly supports academic freedom and a researcher's right to publish; as an institution, however, the Laboratory does not endorse the viewpoint of a publication or guarantee its technical correctness.

SCORR - Supercritical Carbon Dioxide Resist Removal

Gunilla B. Jacobson , Laurie Williams, W. Kirk Hollis, Jerome Barton, Craig M. V. Taylor, Los Alamos National Laboratory, Chemistry Division C-ACT, MS E537, Los Alamos, NM 87545

SCORR, short for supercritical carbon dioxide resist removal, is a new technology that could continue to enable technological development the photolithography processes in industry. SCORR is based upon the physical properties of supercritical fluids (SCFs). These special properties enable SCFs to remove coatings, residues, and particles from high-aspect-ratio structures in integrated circuits (ICs). SCORR also eliminates rinsing and drying steps presently used in IC manufacture. thereby eliminating the generation of millions of gallons of water per fab per day.

Fabricating integrated circuits relies heavily on photolithography to define the shape and pattern of individual components. Once a single stage of a silicon wafer's topography has been completed, the hardened resist must be removed. Conventional processes generates more waste than any single step in the IC manufacturing process, and the production of a complete IC can involve many photolithography iterations. The cost associated with the treatment and disposal of this waste, as well as employee health and safety considerations, are driving a search for environmentally alternative. benign. cost-effective solutions.

In addition, photoresist stripping is confronting finer architectures and higher aspect ratios, as well as new lowk materials that are highly sensitive to

post-etch residue. Low-k dielectrics and low-resistivity conductors such copper are necessary for meeting industry's need for faster and smaller chips. Further, each low-k choice requires different plasma-etching processes. chemistries, or to etch structures into the low-k material: therefore, the nature of the residues can be different. No one product can meet all copper/low-k applications, and existing chemistries are not tunable—or even desirable—for the new processes.

We have developed a new process— SCORR—that known as removes photoresist and post-ash, -etch, and -CMP (particulate) residue semiconductor wafers. As IC feature sizes become smaller, the need for ensuring particle removal will increase. With feature sizes of less than 0.18µm, it will become imperative that all particles greater than about 0.1 micron be removed from the semiconductor wafer. Existing cleaning technologies (such as liquid or high-pressure jet scrubbing) cannot remove particles on the order of 0.1 micron because of surface boundary layer constraints. Because of the low viscosities of supercritical fluids (SCFs). these constraints are virtually eliminated.

The key to the effectiveness of SCORR is the use of SFCs, specifically supercritical carbon dioxide (CO₂). The physical properties (*e.g.*, density, viscosity, diffusivity) of a supercritical fluid often lie between those of gases

and liquids, and can be varied by small changes in temperature and/or pressure.¹

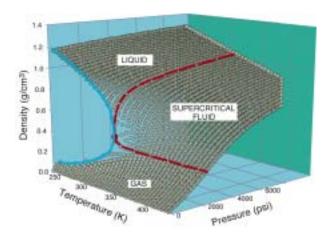


Fig. 1. Carbon dioxide density as a function of temperature and pressure.

The possibility to obtain a wide range of physical properties without changing the chemical composition and by using a fluid than becomes gaseous at atmospheric pressure, makes these fluid attractive for a wide range applications.² Carbon dioxide is generally chemically inert with respect to inorganic materials and is therefore compatible with existing semiconductor materials and metallization systems. Supercritical fluids have, by definition, no surface tension which offers the possibility for the removal of very small (sub-micron) particles from substrate surfaces.³ They are therefore ideal candidates for the removal of photoresist from fabricated surface structures having high aspect ratios. Carbon dioxide is noncombustible, readily available in high purity, environmentally friendly (not ozone depleting), and has a low human toxicity.

Pure supercritical CO₂ is limited in its ability to dissolve most polymers (photoresists).⁴ Rather, it is the

absorption of supercritical CO₂ into the polymer and the resulting swelling of the polymer that can be utilized to achieve various desirable interactions. "tuning" the CO₂/polymer interaction pressure and temperature with adjustments, supercritical CO_2 polymer-based absorbed into the photoresist, thus plasticizing the polymer and lowering its glass transition temperature significantly below that observed at atmospheric pressure. Lowering the polymer's glass transition temperature softens thereby it. enhancing penetration and speeding the diffusion of the supercritical fluid into the polymer/substrate interface.⁵ The principle is to swell the polymer with the dense gas, allowing the supercritical CO₂ to diffuse throughout the polymer matrix and then suddenly release the pressure, forcing a rapid volume change that ruptures the polymer from the substrate.⁶

To further enhance solubility or overcome bonding effects, a second compound, or co-solvent, is added to the supercritical CO₂. The high densities achievable in the SCF region provide the necessary solvating power to solubilize small amounts of co-solvent (less than 5% by volume), which then can diffuse throughout the polymer matrix and to the polymer/substrate interface. This diffusion provides for significant chemical interaction, which promotes the photoresist and residue removal.

Supercritical CO₂ is also attractive from the viewpoint of waste minimization. After the cleaning step, CO₂ is easily separated from the extracted or dissolved compounds, resulting in streams of concentrated compounds and pure, recyclable CO₂. Because SCORR uses pure CO₂ for its

final rinse step, it does not deplete critical water reserves. SCORR's solvent system is not only nontoxic but also recyclable, and thus the only waste generated is the spent solubilized photoresist, which is itself nonregulated. And finally, instead of using alcohol for the drying process, we lower the pressure merely temperature of the superctricial CO₂, and thus the supercritical fluid returns to its gaseous phase, leaving the silicon wafer dry and virtually free of any residue.

The SCORR system technology based on sound green chemistry principles while enabling the industry to meet its development goals, such as the introduction of low-k dielectric materials, which are sensitive to the established chemistries, plus copper and smaller line widths. The chemistry SCORR system instrumentation will be presented in detail. Advantages of the SCORR system to the industry and the environment will also be discussed.

¹ a) Squires, T. G.; Venier, C. G.; Aida, T. *Fluid Phase Equilibria*, **1983**, 10, 261-268. b) Tiltscher, H.; Hofmann, H. *Chem. Eng. Sci.***1987**, 42, 959-977.

² Clifford, T. Fundamentals of Supercritical Fluids, Oxford University Press, Inc., New York, 1999.

³ Manivan, G.; Sawan, S. P., Supercritical Fluid Cleaning: Fundamentals, Technology, and Applications, McHardy, J., editor, Noyes Publications, Westwood, New Jersey, 1998.

⁴ DeSimone, J. M.; Guan, Z.; Elsbernd, C. S. Science, **1992**, 257, 945-947.

⁵ Chiou, J. S.; Barlow, J. W.; Paul, D. R. *J. Appl. Polym. Sci.*, **1985**, 30, 2633-2642.

⁶ Barton, J. U.S. Patent 6 085 762, 2000.